C,D,E: Plumule formation and adventitious bud elongation

A: Callus induction; B: Adventitious bud recovery;

Fig.1 Adventitious bud induction from leaf

1/11

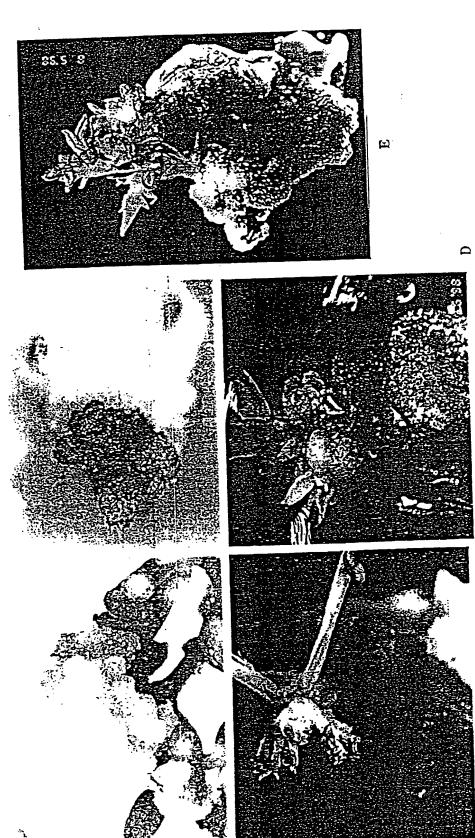


Fig. 2 Acacia mangium regeneration from petiole. A: Callus induction; B: Adventitious bud recovery; C: Plumule formation and adventitious bud elongation.

Ö

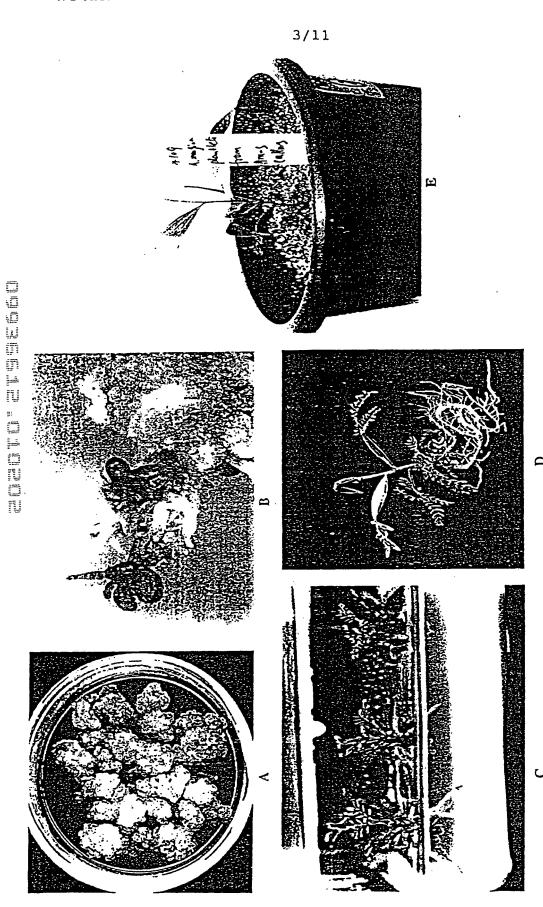
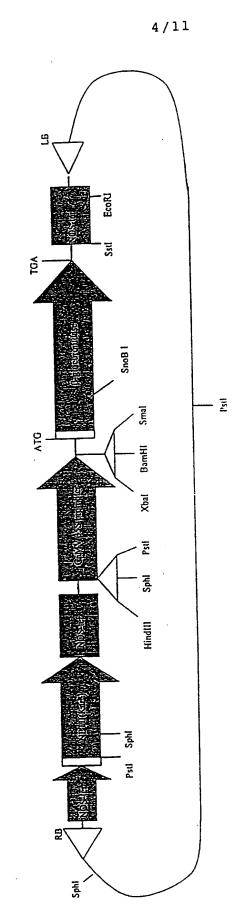


Fig.3 Acacia mangium regeneration from stem pieces A: Callus; B: Adventitious buds induction from callus; C: Root formation; D: Complete plantlet; E: One month old plantlet in pot soil.

pB1121 vector



mosaic virus (CaMV) 35S promoter cloned upstream of the GUS gene. Vector size: 13.0kb(CLONTECH) Fig.4 Map of pBI121 with an 800-bp HindIII-BamHI fragment containing the cauliflower

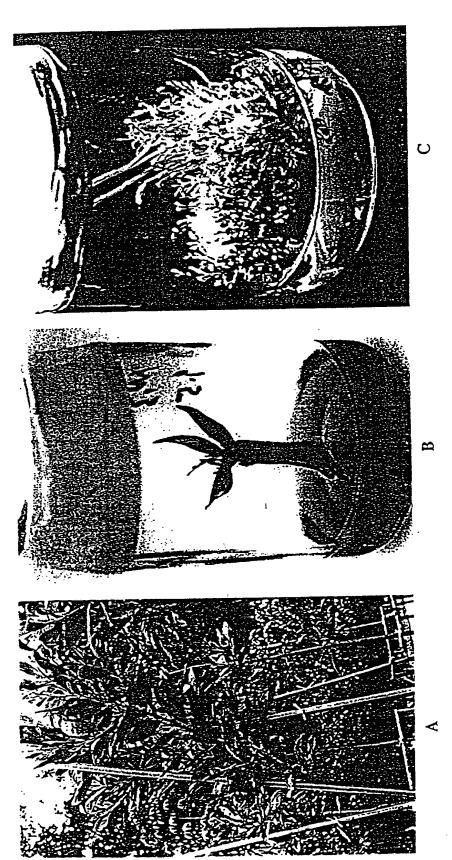


Fig.5 Rejuvenation of tree. A: Two-year old tree; B: Adventitious bud induction; C: Propagated adventitious buds with plumules

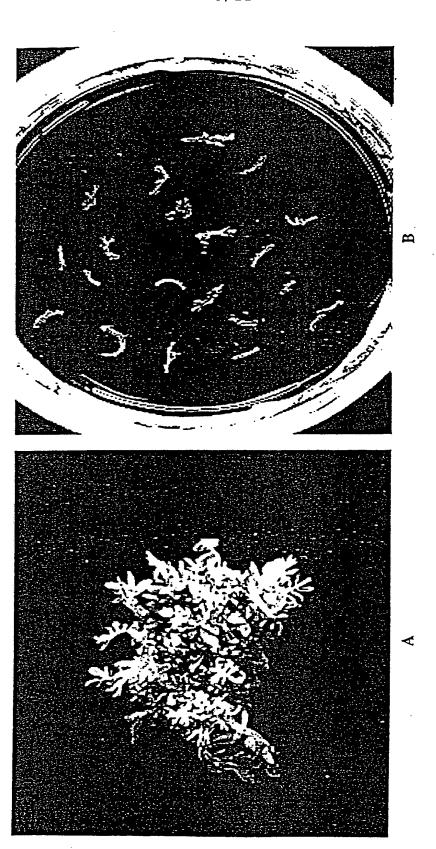


Fig.6 A: Adventitious buds; B: Stem pieces as explants for transformation

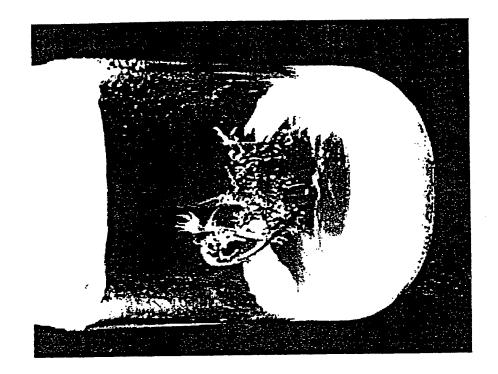


Fig. 7 Selection and induction of putative transgenic adventitious buds

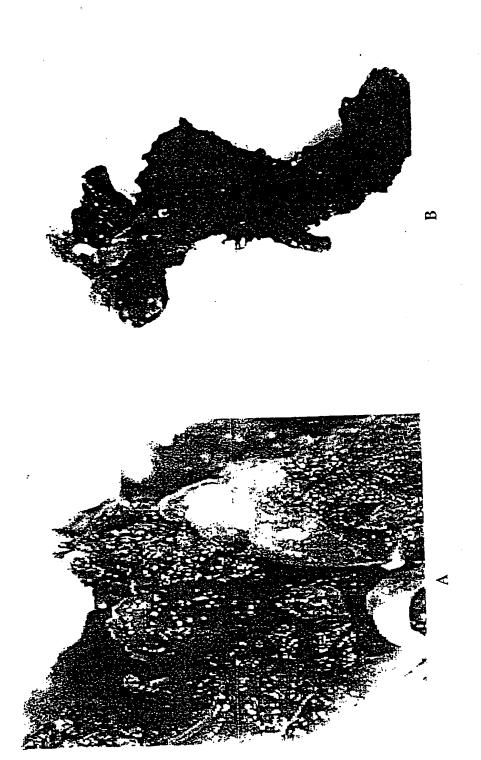


Fig. 8 Gus staining of adventitious buds after selection for 5 months





Fig.9 GUS staining of young transgenic stem pieces
A,B: stem pieces
C: A shoot



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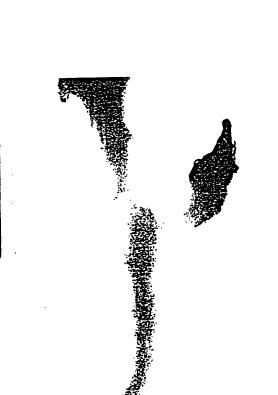


Fig.10 GUS staining of transgenic leaf and leaf pieces '

5 4 3 ck 2 1

—— 23 kb —— 9.4 kb —— 6.5 kb —— 6.5 kb --- 2.3 kb

from non-transgenic plant; 7, positive control: DNA from a tomato transgenic line Fig.11 Southern blot to nptll probe: 20 µg DNA was digested with Hind III and hybridized to nptII probe. 1-6, transgenic lines; ck, negative control: DNA by plasmid pWS42 with nptII as selection maker.

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